WHAT IS CLAIMED IS:

1		1.	A method for loading a disaccharide into mammalian nucleated cells,			
2	comprising:					
3		conta	cting said cells for at least 2 hours with a solution comprising at least one			
4	disaccharide,	e, thereby loading the cells with disaccharide to produce disaccharide-loaded				
5	mammalian n	nmalian nucleated cells.				
1		2.	A method of claim 1, wherein said cells are selected from the group			
2	consisting of	nsisting of stem cells, immune system cells, and epithelial cells.				
1		3.	A method of claim 1, wherein said contacting is for 10 hours.			
1		4.	A method of claim 1, wherein said contacting is for 24 hours.			
1		5.	A method of claim 1, wherein said disaccharide is trehalose.			
1		6.	A method of claim 1, wherein said solution further comprises not more			
2	than 3% dimethyl sulfoxide.					
1		7.	A method for increasing survival of mammalian nucleated cells			
2	following dry	ying and rehydration, comprising:				
3		(a) contacting said cells with a solution comprising at least one disaccharide				
4	for at least 21	or at least 2 hours, thereby producing disaccharide-loaded cells,				
5		(b) drying said disaccharide-loaded cells to a residual water content between				
6	0.2 and 0.5 gr	0.2 and 0.5 gram water per gram of dry weight, and				
7		(c) rehydrating said cells,				
8	thereby increasing survival of the cells.					
1		8.	A method of claim 7, wherein said contacting is for 24 hours.			
1		9.	A method of claim 7, wherein said cells are selected from the group			
2	consisting of stem cells, immune system cells, and epithelial cells.					
1		10.	A method of claim 7, wherein said disaccharide is trehalose.			
1		11.	A method of claim 7, wherein said cells further comprise a heat shock			
2	protein.					

1 12. A method of claim 11, wherein said heat shock protein is induced by 2 exposing said cells to a heat shock.

- 1 13. A method of claim 12, wherein said heat shock consists of raising the temperature of medium contacting the cells to 42 44 °C for one hour, and then allowing the temperature of the medium to drop to 36-38 °C.
- 1 14. A method of claim 11, wherein said heat shock protein is introduced 2 into the cells by contacting said cells with a solution comprising said protein.
- 1 15. A method of claim 11, wherein said heat shock protein is expressed 2 from a nucleic acid sequence introduced into said cells.
- 1 16. A method of claim 11, wherein said heat shock protein is p26 from 2 Artemia franciscana.
- 1 17. A method of claim 7, further wherein said cells are contacted with a solution comprising an apoptosis inhibitor.
- 1 18. A method of claim 17, wherein said apoptosis inhibitor is selected
- 2 from the group consisting of N-(2-Quinolyl)valyl-asparty1-(2,6-difluorophenoxy)methyl
- 3 ketone (in which the aspartyl residue is o-methylated or non-o-methylated), caspase I
- 4 inhibitor II, calpain inhibitor, and Bcl-xL.
- 1 19. A method of claim 7, further wherein said cells are contacted by a solution comprising arbutin or hydroquinone, provided that said cells are not 293 cells or B cells.
- 1 20. A method of claim 7, further wherein said cells are contacted by a solution comprising not more than 3% dimethyl sulfoxide.
- 1 21. A method of claim 7, further wherein said cells are contacted by a solution comprising a heat shock protein and an apoptosis inhibitor.
- 1 22. A method of claim 21, wherein said solution further comprises not 2 more than 3% dimethyl sulfoxide.

1	23. A method of claim 19, wherein said cells are dried in a medium					
2	comprising arbutin or hydroquinone.					
1	24. A method of claim 7, wherein said cells are dried in rounded droplets					
2	of drying buffer.					
1	25. A method for increasing survival of mammalian nucleated cells					
2	following drying and rehydration, comprising:					
3	(a) contacting said cells with a solution comprising an apoptosis inhibitor,					
4	thereby loading the cells with said apoptosis inhibitor, to produce apoptosis inhibitor -loaded					
5	cells,					
6	(b) drying said apoptosis inhibitor-loaded cells, and					
7	(c) rehydrating said cells,					
8	thereby increasing survival of the cells.					
1	26. A method of claim 25, wherein said apoptosis inhibitor is selected					
2	from the group consisting of N-(2-Quinolyl)valyl-aspartyl-(2,6-difluorophenoxy)methyl					
3	ketone (in which the aspartyl residue is o-methylated or non-o-methylated), Caspase I					
4	inhibitor II, Calpain inhibitor, and Bcl-xL.					
1	27. A method of claim 25, wherein said cells are selected from the group					
2	consisting of stem cells, immune system cells, and epithelial cells					
1	28. A method of claim 25, wherein said cells are dried in droplets of					
2	drying buffer.					
1	29. A method for increasing survival of mammalian nucleated cells					
2	following drying and rehydration, comprising:					
3	(a) introducing a heat shock protein into, or inducing production of a heat					
4	shock protein in, said cells, to produce heat shock protein-loaded cells,					
5	(b) drying said heat shock protein-loaded cells, and					
5	(c) rehydrating said cells,					
7	thereby increasing survival of the cells.					
1	30. A method of claim 29, wherein said heat shock protein is p26 from					

Artemia franciscana.

1	31. A method of claim 29, wherein said heat shock protein is introduced				
2	into said cells by incubating said cells in a medium comprising said heat shock protein.				
1	32. A method of claim 29, wherein said heat shock protein is induced in				
2	said cells by raising the temperature of medium contacting the cells to 42 - 44 °C for one				
3	hour, and then allowing the temperature of the medium to lower to 36-38 °C.				
1	33. A method of claim 29, wherein said heat shock protein is introduced				
2	into said cells by introducing into said cells a nucleic acid sequence comprising a promoter				
3	operably linked to a sequence encoding said heat shock protein.				
1	34. A method of claim 29, wherein said cells are selected from the group				
2	consisting of stem cells, immune system cells, and epithelial cells.				
1	35. A method of claim 29, wherein said cells are dried in droplets of				
2	drying buffer.				
1	36. A method for increasing survival of mammalian nucleated cells				
2	following drying and rehydration, provided said cells are not 293 cells or B cells, comprising				
3	(a) incubating said cells with a compound selected from arbutin and				
4	hydroquinone, to produce arbutin- or hydroquinone- loaded cells,				
5	(b) drying said arbutin- or hydroquinone- loaded cells, and				
6	(c) rehydrating said cells,				
7	thereby increasing survival of the cells.				
1	37. A method of claim 36, wherein said compound of step (a) is arbutin.				
1	38. An isolated mammalian nucleated cell comprising a disaccharide and				
2	compound selected from the group consisting of arbutin and hydroquinone.				
1	39. An isolated mammalian nucleated cell of claim 38, wherein said				
2	compound is arbutin.				
1	40. A mammalian nucleated cell of claim 38, wherein said cell is dried.				
1	41. A mammalian nucleated cell of claim 38, further comprising an				
2	apoptosis inhibitor.				

1		42.	A mammalian nucleated cell of claim 38, further comprising a heat		
2	shock protein	•			
1		43.	A mammalian nucleated cell of claim 38, wherein said disaccharide is		
2	trehalose.				
1		44.	An isolated dried mammalian nucleated cell comprising a disaccharide		
2	and an exogenous heat shock protein.				
1		45.	A dried mammalian nucleated cell of claim 44, wherein said		
2	disaccharide is trehalose.				
1		46.	A isolated, dried mammalian nucleated cell comprising a disaccharide		
2	and an exogenous apoptosis inhibitor.				
1		47.	A dried mammalian nucleated cell of claim 46, wherein said		
2	disaccharide is trehalose.				